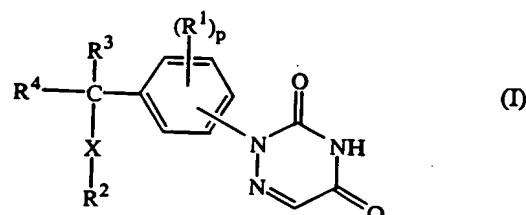


ABSTRACT

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IL-5 INHIBITING 6-AZAUURACIL DERIVATIVES

The present invention is concerned with the compounds of formula



the *N*-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein p is 0 to 4; X is O, S, NR⁵ or a direct bond; Y is O, S, NR⁵ or S(O)₂; R¹ independently is C₁-6alkyl, halo, polyhaloC₁-6alkyl, hydroxy, mercapto, C₁-6alkyloxy, C₁-6alkylthio, C₁-6alkylcarbonyloxy, aryl, cyano, nitro, Het³, R⁶, NR⁷R⁸ or substituted C₁-4alkyl; R² is Het¹, C₃-7cycloalkyl or optionally substituted C₁-6alkyl and if X is O, S or NR⁵, then R² may also represent aminocarbonyl, 15 aminothiocarbonyl, C₁-4alkylcarbonyl, C₁-4alkylthiocarbonyl, arylcarbonyl, arylthiocarbonyl, Het¹carbonyl or Het¹thiocarbonyl; R³ and R⁴ independently are hydrogen, C₁-6alkyl or C₃-7cycloalkyl; R³ and R⁴ form a C₂-6alkanediyl; R⁵ is hydrogen or C₁-4alkyl; R⁶ is a sulfonyl or sulfinyl derivative; R⁷ and R⁸ are independently hydrogen, optionally substituted C₁-4alkyl, aryl, a carbonyl containing moiety, C₃-7cycloalkyl, 20 -Y-C₁-4alkanediyl-C(=O)-O-R¹⁴, Het³, Het⁴ and R⁶; R¹¹ is hydroxy, mercapto, cyano, nitro, halo, trihalomethyl, C₁-4alkyloxy, formyl, trihaloC₁-4alkylsulfonyloxy, R⁶, NR⁷R⁸, C(=O)NR⁷R⁸, C₁-4alkanediyl-C(=O)-O-R¹⁴, -C(=O)-O-R¹⁴, -Y-C₁-4alkanediyl-C(=O)-O-R¹⁴, aryl, aryloxy, arylcarbonyl, C₃-7cycloalkyl, C₃-7cycloalkyloxy, phthalimide-2-yl, Het³ and C(=O)Het³; R¹⁴ is hydrogen, C₁-4alkyl, C₃-7cycloalkyl, aminocarbonylmethylene or 25 mono- or di(C₁-4alkyl)aminocarbonylmethylene; aryl is optionally substituted phenyl; Het¹, Het², Het³ and Het⁴ are optionally substituted heterocycles; to processes for their preparation and compositions comprising them. It further relates to their use as a medicine.